

L Number	Hits	Search Text	DB	Time stamp
1	813	(proteons or PNC or "misfolded proteins")	USPAT; US-PGPUB	2004/07/27 14:22
2	934	(proteons or PNC or "misfolded proteins")	USPAT; US-PGPUB; EPO; JPO; DERWENT	2004/07/27 14:22
3	11437	heating and centrifuging	USPAT; US-PGPUB; EPO; JPO; DERWENT	2004/07/27 14:23
4	10	(heating and centrifuging) and ((proteons or PNC or "misfolded proteins"))	USPAT; US-PGPUB; EPO; JPO; DERWENT	2004/07/27 14:23

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FULL ESTIMATED COST	0.21	0.21

=> file Uspatfull

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.42	0.42

FILE 'USPATFULL' ENTERED AT 15:45:47 ON 27 JUL 2004

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FILE COVERS 1971 TO PATENT PUBLICATION DATE: 27 Jul 2004 (20040727/PD)

FILE LAST UPDATED: 27 Jul 2004 (20040727/ED)

HIGHEST GRANTED PATENT NUMBER: US6769133

HIGHEST APPLICATION PUBLICATION NUMBER: US2004143878

CA INDEXING IS CURRENT THROUGH 27 Jul 2004 (20040727/UPCA)

ISSUE CLASS FIELDS (/INCL) CURRENT THROUGH: 27 Jul 2004 (20040727/PD)

REVISED CLASS FIELDS (/NCL) LAST RELOADED: Jun 2004

USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Jun 2004

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This file contains CAS Registry Numbers for easy and accurate
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=> (proteons or prions "misfolded proteins") and (heating and centrifuging)

8 PROTEONS
994 PRIONS
2311 "MISFOLDED"
149454 "PROTEINS"
0 PRIONS "MISFOLDED PROTEINS"
(PRIONS (W) "MISFOLDED" (W) "PROTEINS")
665771 HEATING
558 HEATINGS
665810 HEATING
(HEATING OR HEATINGS)
25206 CENTRIFUGING
24 CENTRIFUGINGS
25220 CENTRIFUGING
(CENTRIFUGING OR CENTRIFUGINGS)
L1 1 (PROTEONS OR PRIONS "MISFOLDED PROTEINS") AND (HEATING AND CENTRIFUGING)

=> (proteons or prions or "misfolded proteins") and (heating and centrifuging)

8 PROTEONS
994 PRIONS
2311 "MISFOLDED"
149454 "PROTEINS"
167 "MISFOLDED PROTEINS"
("MISFOLDED" (W) "PROTEINS")
665771 HEATING
558 HEATINGS
665810 HEATING
(HEATING OR HEATINGS)
25206 CENTRIFUGING
24 CENTRIFUGINGS
25220 CENTRIFUGING
(CENTRIFUGING OR CENTRIFUGINGS)
L2 33 (PROTEONS OR PRIONS OR "MISFOLDED PROTEINS") AND (HEATING AND CENTRIFUGING)

=> d ti 1-33

L2 ANSWER 1 OF 33 USPATFULL on STN
TI Method of isolation and self-assembly of small protein particles from blood and other biological materials

L2 ANSWER 2 OF 33 USPATFULL on STN
TI Multi-formed collagenous biomaterial medical device

L2 ANSWER 3 OF 33 USPATFULL on STN
TI Yeast cell surface display of proteins and uses thereof

L2 ANSWER 4 OF 33 USPATFULL on STN
TI Compositions and methods for detection and isolation of phosphorylated molecules

L2 ANSWER 5 OF 33 USPATFULL on STN
TI Yeast cell surface display of proteins and uses thereof

L2 ANSWER 6 OF 33 USPATFULL on STN
TI Methods for sterilizing biological materials by irradiation over a temperature gradient

L2 ANSWER 7 OF 33 USPATFULL on STN

TI Cell and tissue culture device

L2 ANSWER 8 OF 33 USPATFULL on STN
TI Surrogate antibodies and methods of preparation and use thereof

L2 ANSWER 9 OF 33 USPATFULL on STN
TI Methods and compositions for neutralizing anthrax and other bioagents

L2 ANSWER 10 OF 33 USPATFULL on STN
TI Radiopaque implantable collagenous biomaterial device

L2 ANSWER 11 OF 33 USPATFULL on STN
TI Method and system for detecting and recording submicron sized particles

L2 ANSWER 12 OF 33 USPATFULL on STN
TI Methods for sterilizing tissue

L2 ANSWER 13 OF 33 USPATFULL on STN
TI Methods for sterilizing tissue

L2 ANSWER 14 OF 33 USPATFULL on STN
TI Recombinant collagen and derived proteins produced by plants, methods for obtaining them and uses

L2 ANSWER 15 OF 33 USPATFULL on STN
TI Targeted enzyme prodrug therapy

L2 ANSWER 16 OF 33 USPATFULL on STN
TI Recombinant collagens and derived proteins produced by plants, methods of obtaining such and their uses

L2 ANSWER 17 OF 33 USPATFULL on STN
TI Antibodies for specifically detecting pathogenic prions of human origin, and detection methods carried out using these antibodies

L2 ANSWER 18 OF 33 USPATFULL on STN
TI Targeted enzymes

L2 ANSWER 19 OF 33 USPATFULL on STN
TI Tubular graft construct

L2 ANSWER 20 OF 33 USPATFULL on STN
TI Adjuvants for nucleic acid vaccines

L2 ANSWER 21 OF 33 USPATFULL on STN
TI Methods devices and systems for sorting and separating particles

L2 ANSWER 22 OF 33 USPATFULL on STN
TI Rapid sterilization and vaccine preparation

L2 ANSWER 23 OF 33 USPATFULL on STN
TI Nucleic acid sequences encoding type III tenebrio antifreeze proteins and method for assaying activity

L2 ANSWER 24 OF 33 USPATFULL on STN
TI Nucleic acid sequences encoding type III tenebrio antifreeze proteins and method for assaying activity

L2 ANSWER 25 OF 33 USPATFULL on STN

TI MULTI-FORMED COLLAGENOUS BIOMATERIAL MEDICAL DEVICE

L2 ANSWER 26 OF 33 USPATFULL on STN
TI Yeast cell surface display of proteins and uses thereof

L2 ANSWER 27 OF 33 USPATFULL on STN
TI Elemental analysis of tagged biologically active materials

L2 ANSWER 28 OF 33 USPATFULL on STN
TI Assay for specific strains of multiple disease related conformations of a protein

L2 ANSWER 29 OF 33 USPATFULL on STN
TI Yeast cell surface display of proteins and uses thereof

L2 ANSWER 30 OF 33 USPATFULL on STN
TI Nucleic acid-coupled colorimetric analyte detectors

L2 ANSWER 31 OF 33 USPATFULL on STN
TI Yeast cell surface display of proteins and uses thereof

L2 ANSWER 32 OF 33 USPATFULL on STN
TI Microbial inactivation by high-pressure throttling

L2 ANSWER 33 OF 33 USPATFULL on STN
TI Production of insulin-like growth factor-1 in methylotrophic yeast cells

=> d ab bib 2, 6, 17, 28

L2 ANSWER 2 OF 33 USPATFULL on STN
AB The invention involves a submucosa tissue that has the capability of being shape formed or shape configured. The submucosa involves a purified form of submucosa tissue. Optionally, the submucosa can be packaged in such a manner to permit sterility or maintain sterility of the submucosa.

AN 2004:177878 USPATFULL
TI Multi-formed collagenous biomaterial medical device
IN Hiles, Michael C., Lafayette, IN, UNITED STATES
Patel, Umesh H., West Lafayette, IN, UNITED STATES

PI US 2004137042 A1 20040715
AI US 2003-744420 A1 20031223 (10)

RLI Continuation of Ser. No. US 1999-320933, filed on 27 May 1999, GRANTED, Pat. No. US 6666892 Continuation-in-part of Ser. No. US 1997-916490, filed on 22 Aug 1997, GRANTED, Pat. No. US 6206931

PRAI US 1996-24542P 19960823 (60)
US 1996-24693P 19960906 (60)
US 1998-110407P 19981201 (60)

DT Utility
FS APPLICATION

LREP Woodard, Emhardt, Moriarty, McNett & Henry LLP, Bank One Center/Tower, Suite 3700, 111 Monument Circle, Indianapolis, IN, 46204-5137

CLMN Number of Claims: 20
ECL Exemplary Claim: 1
DRWN 1 Drawing Page(s)
LN.CNT 937

L2 ANSWER 6 OF 33 USPATFULL on STN
AB Methods are disclosed for sterilizing tissue to reduce the level of one

or more active biological contaminants or pathogens therein, such as viruses, bacteria, (including inter- and intracellular bacteria, such as mycoplasmas, ureaplasmas, nanobacteria, chlamydia, rickettsias), yeasts, molds, fungi, spores, **prions** or similar agents responsible, alone or in combination, for TSEs and/or single or multicellular parasites. The methods involve sterilizing one or more tissues with irradiation.

AN 2004:44170 USPATFULL

TI Methods for sterilizing biological materials by irradiation over a temperature gradient

IN MacPhee, Martin, Montgomery Village, MD, UNITED STATES

Calvert, Glenn, Frederick, MD, UNITED STATES

Lynch, Tom, Rockville, MD, UNITED STATES

Kent, Randall S., Thousand Oaks, CA, UNITED STATES

Mann, David, Gaithersburg, MD, UNITED STATES

Burgess, Wilson, Clifton, VA, UNITED STATES

Miekka, Shirley, Gaithersburg, MD, UNITED STATES

Drohan, William N., Springfield, VA, UNITED STATES

PI US 2004033160 A1 20040219

AI US 2002-197249 A1 20020718 (10)

DT Utility

FS APPLICATION

LREP FLESHNER & KIM, LLP, P.O. BOX 221200, CHANTILLY, VA, 20153

CLMN Number of Claims: 102

ECL Exemplary Claim: 1

DRWN 2 Drawing Page(s)

LN.CNT 3969

L2 ANSWER 17 OF 33 USPATFULL on STN

AB Antibodies for specifically detecting pathogenic **prions** of human origin, and methods for detecting pathogenic **prions**, are described. In particular, a conformation-dependent immunoassay method for detecting pathogenic prion proteins in a sample of a body fluid, containing a PrP protein, which contains a first, natural, non-pathological conformation, i.e. PrP.sup.c, and a second, pathological conformation, i.e. PrP.sup.Sc, is described, in which method the prion proteins differ in their binding affinity for monoclonal antibodies which bind specifically to prion proteins of human origin, with the detection method comprising the following steps:

a) adding one of the abovementioned monoclonal antibodies, which is fixed to a solid support and which exhibits a higher affinity for the first prion protein conformation, to the first portion of the sample, and determining this first concentration;

b) treating the second portion of the sample in order to increase the binding affinity of the second conformation of the prion protein for the monoclonal antibody;

c) adding the monoclonal antibody to the treated second portion of the sample to be investigated, in order to determine the second concentration;

d) comparing the first prion protein concentration with the second prion protein concentration in order to ascertain the presence of the pathogenic prion protein conformation.

AN 2003:134009 USPATFULL

TI Antibodies for specifically detecting pathogenic **prions** of human origin, and detection methods carried out using these antibodies

IN Vey, Martin, Marburg, GERMANY, FEDERAL REPUBLIC OF
Lang, Wiegand, Coelbe, GERMANY, FEDERAL REPUBLIC OF
Groener, Albrecht, Marburg, GERMANY, FEDERAL REPUBLIC OF
Bellon, Anne, Marburg, GERMANY, FEDERAL REPUBLIC OF
PI US 2003092094 A1 20030515
AI US 2002-273282 A1 20021018 (10)
PRAI DE 2001-152677 20011019
DT Utility
FS APPLICATION
LREP Finnegan, Henderson, Farabow,, Garrett & Dunner, L.L.P., 1300 I Street,
N.W., Washington, DC, 20005-3315
CLMN Number of Claims: 22
ECL Exemplary Claim: 1
DRWN 3 Drawing Page(s)
LN.CNT 708
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L2 ANSWER 28 OF 33 USPATFULL on STN
AB Assay methodology of the invention allows for: (1) determining if a
sample contains a conformation of a protein which is associated with
disease and the concentration and amount of such if present; (2)
determining the amount of protease resistant disease related protein in
a sample and by subtracting that amount from the total amount of disease
related protein present determining the amount of protease sensitive
disease protein in the sample; and (3) determining the strain and
incubation time of a disease related protein by (i) relating the
relative amounts of protease resistant and protease sensitive protein to
known strains to thereby determine the strain; and (ii) plotting the
concentration of protease sensitive protein on a graph of incubation
time versus concentration of protease sensitive protein for known
strains to predict the incubation time of an unknown strain of
pathogenic protein in a sample.
AN 2002:3842 USPATFULL
TI Assay for specific strains of multiple disease related conformations of
a protein
IN Prusiner, Stanley B., San Francisco, CA, UNITED STATES
Safar, Jiri G., Concord, CA, UNITED STATES
Cohen, Fred E., San Francisco, CA, UNITED STATES
PI US 2002001817 A1 20020103
US 6617119 B2 20030909
AI US 2001-901865 A1 20010709 (9)
RLI Continuation of Ser. No. US 1998-151057, filed on 10 Sep 1998, PENDING
Continuation-in-part of Ser. No. US 1998-26957, filed on 20 Feb 1998,
ABANDONED Continuation-in-part of Ser. No. US 1997-804536, filed on 21
Feb 1997, GRANTED, Pat. No. US 5891641
DT Utility
FS APPLICATION
LREP Karl Bozicevic, Bozicevic, Field and Francis LLP, Suite 200, 200
Middlefield Road, Menlo Park, CA, 94025
CLMN Number of Claims: 20
ECL Exemplary Claim: 1
DRWN 19 Drawing Page(s)
LN.CNT 2676
CAS INDEXING IS AVAILABLE FOR THIS PATE

27/07/200411:38Print selected from Online session

=> index bioscience patents

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FILE 'ENCOMPPAT' ACCESS NOT AUTHORIZED

FILE 'ENCOMPPAT2' ACCESS NOT AUTHORIZED

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SESSION

FULL ESTIMATED COST

0.21

0.21

INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, AQUASCI, BIOBUSINESS, BIOCOMMERCE, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA, CANCERLIT, CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB, CROPU, DISSABS, DDFB, DDFU, DGENE, DRUGB, DRUGMONOG2, ...' ENTERED AT 11:02:29 ON 27 JUL 2004

92 FILES IN THE FILE LIST IN STNINDEX

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=> "cyclic amplification" and "proteons"

1 FILE DGENE

27 FILES SEARCHED...

1 FILE IFIPAT

60 FILES SEARCHED...

1 FILE USPATFULL

83 FILES SEARCHED...

1 FILE PCTFULL

4 FILES HAVE ONE OR MORE ANSWERS, 92 FILES SEARCHED IN STNINDEX

L1 QUE "CYCLIC AMPLIFICATION" AND "PROTEONS"

=> file dgebe ifipat uspatfull pctfull

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4.20

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=> "cyclic amplification" and "proteons"

L2 4 "CYCLIC AMPLIFICATION" AND "PROTEONS"

=> d ti 1-4

27/07/200411:38Print selected from Online session

L2 ANSWER 1 OF 4 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN
TI Cyclically amplifying **proteons** in a biological sample for
detecting conformational or congophilic disorders by centrifuging the
sample until a supernatant is formed, dividing the supernatant into
subsamples and heating a subsample.

L2 ANSWER 2 OF 4 IFIPAT COPYRIGHT 2004 IFI on STN
TI METHOD OF ISOLATION AND SELF-ASSEMBLY OF SMALL PROTEIN PARTICLES FROM
BLOOD AND OTHER BIOLOGICAL MATERIALS

L2 ANSWER 3 OF 4 USPATFULL on STN
TI Method of isolation and self-assembly of small protein particles from
blood and other biological materials

L2 ANSWER 4 OF 4 PCTFULL COPYRIGHT 2004 Univentio on STN
TIEN METHOD OF ISOLATION AND SELF-ASSEMBLY OF SMALL PROTEIN PARTICLES FROM
BLOOD AND OTHER BIOLOGICAL MATERIALS
TIFR METHODES D'ISOLEMENT ET D'AUTO-ASSEMBLAGE DE PETITES PARTICULES
PROTEINIQUES PRESENTES DANS DU SANG OU DANS D'AUTRES MATIERES
BIOLOGIQUES

=> index bioscience patents

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FILE 'ENCOMPPAT' ACCESS NOT AUTHORIZED

FILE 'ENCOMPPAT2' ACCESS NOT AUTHORIZED

COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
4.97	9.17

FULL ESTIMATED COST

INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, AQUASCI, BIOBUSINESS,
BIOCOMMERCE, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA, CANCERLIT,
CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB, CROPU, DISSABS, DDFB, DDFU,
DGENE, DRUGB, DRUGMONOG2, ...' ENTERED AT 11:07:30 ON 27 JUL 2004

92 FILES IN THE FILE LIST IN STNINDEX

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=> ("cyclic amplification" or amplification) and ("misfolded proteins" or proteins
or PNC)

3	FILE ADISCTI
3	FILE ADISINSIGHT
1	FILE ADISNEWS
413	FILE AGRICOLA
29	FILE ANABSTR
135	FILE AQUASCI
249	FILE BIOBUSINESS
12	FILE BIOCOMMERCE
6054	FILE BIOSIS
4371	FILE BIOTECHABS
4371	FILE BIOTECHDS
2953	FILE BIOTECHNO
1155	FILE CABA
4800	FILE CANCERLIT
10613	FILE CAPLUS
53	FILE CEABA-VTB
29	FILE CEN

18 FILE CIN
4 FILE CONFSCI
19 FILES SEARCHED...
14 FILE CROPU
427 FILE DISSABS
1 FILE DDFB
22 FILE DDFU
101731 FILE DGENE
1 FILE DRUGB
3 FILE IMSDRUGNEWS
82 FILE DRUGU
2 FILE IMSRESEARCH
30 FILES SEARCHED...
33 FILE EMBAL
3283 FILE EMBASE
2528 FILE ESBIODBASE
327 FILE FEDRIP
49 FILE FROSTI
82 FILE FSTA
5791 FILE GENBANK
1 FILE HEALSAFE
1669 FILE IFIPAT
123 FILE JICST-EPLUS
8 FILE KOSMET
1813 FILE LIFESCI
10820 FILE MEDLINE
6 FILE NIOSHTIC
95 FILE NTIS
32 FILE OCEAN
51 FILES SEARCHED...
1171 FILE PASCAL
1 FILE PHAR
4 FILE PHARMAML
55 FILE PHIN
432 FILE PROMT
11 FILE RDISCLOSURE
4749 FILE SCISEARCH
2384 FILE TOXCENTER
45430 FILE USPATFULL
2134 FILE USPAT2
12 FILE VETU
1493 FILE WPIDS
9 FILE WPIFV
1493 FILE WPINDEX
1 FILE CAOLD
1 FILE CASREACT
72 FILES SEARCHED...
28 FILE DPCI
6830 FILE EUROPATFULL
216 FILE FRFULL
201 FILE INPADOC
5 FILE JAPIO
18 FILE PAPERCHEM2
304 FILE PATDPAFULL
61 FILE PATOSEP
79 FILE PATOSWO
32591 FILE PCTFULL
87 FILES SEARCHED...
3 FILE RAPRA

71 FILES HAVE ONE OR MORE ANSWERS, 92 FILES SEARCHED IN STNINDEX

L3 QUE ("CYCLIC AMPLIFICATION" OR AMPLIFICATION) AND ("MISFOLDED PROTEINS" OR PROTEINS OR PNC)

=> "cyclic amplification" and ("misfolded proteins" or PNC)

1 FILE BIOSIS
1 FILE BIOTECHNO
1 FILE CAPLUS

24 FILES SEARCHED...

1 FILE DGENE
1 FILE EMBASE
1 FILE ESBIODBASE
1 FILE IFIPAT
1 FILE LIFESCI

45 FILES SEARCHED...

1 FILE MEDLINE
1 FILE SCISEARCH
1 FILE USPATFULL

70 FILES SEARCHED...

1 FILE PCTFULL

90 FILES SEARCHED...

12 FILES HAVE ONE OR MORE ANSWERS, 92 FILES SEARCHED IN STNINDEX

L4 QUE "CYCLIC AMPLIFICATION" AND ("MISFOLDED PROTEINS" OR PNC)

=> file biosis biotechno caplus embase esbiobase lifesci medline scisearch
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FULL ESTIMATED COST	7.98	17.15

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=> "cyclic amplification" and ("misfolded proteins" or PNC)

L5 8 "CYCLIC AMPLIFICATION" AND ("MISFOLDED PROTEINS" OR PNC)

=> d ti 1-8

L5 ANSWER 1 OF 8 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN

TI **Cyclic amplification** of protein misfolding:
Application to prion-related disorders and beyond.

L5 ANSWER 2 OF 8 BIOTECHNO COPYRIGHT 2004 Elsevier Science B.V. on STN

TI **Cyclic amplification** of protein misfolding:
Application to prion-related disorders and beyond

L5 ANSWER 3 OF 8 CAPLUS COPYRIGHT 2004 ACS on STN

TI **Cyclic amplification** of protein misfolding:
application to prion-related disorders and beyond

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TI **Cyclic amplification** of protein misfolding:
Application to prion-related disorders and beyond.

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TI **Cyclic amplification** of protein misfolding:
Application to prion-related disorders and beyond

L5 ANSWER 6 OF 8 LIFESCI COPYRIGHT 2004 CSA on STN

TI **Cyclic amplification** of protein misfolding:
application to prion-related disorders and beyond

L5 ANSWER 7 OF 8 MEDLINE on STN

TI **Cyclic amplification** of protein misfolding:
application to prion-related disorders and beyond.

L5 ANSWER 8 OF 8 SCISEARCH COPYRIGHT 2004 THOMSON ISI on STN

TI **Cyclic amplification** of protein misfolding:
application to prion-related disorders and beyond

=> d ab bib 1

L5 ANSWER 1 OF 8 BIOSIS COPYRIGHT 2004 BIOLOGICAL ABSTRACTS INC. on STN

AB Diverse human disorders, including the majority of neurodegenerative diseases, are thought to arise from the misfolding and aggregation of protein. We have recently described a novel technology to amplify cyclically **misfolded proteins** in vitro. This procedure, named protein misfolding **cyclic amplification** (PMCA), is conceptually analogous to DNA amplification by PCR and has tremendous implications for research and diagnosis. The PMCA concept has been proved on the amplification of prions implicated in the pathogenesis of transmissible spongiform encephalopathies. In this article we describe the rational behind PMCA and some of the many potential applications of this novel technology.

AN 2002:430276 BIOSIS

DN PREV200200430276

TI **Cyclic amplification** of protein misfolding:
Application to prion-related disorders and beyond.

AU Soto, Claudio [Reprint author]; Saborio, Gabriela P.; Anderes, Laurence

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CS 15 Chemin des Mines, 1211, Geneva, Switzerland
claudio.soto@serono.com
SO Trends in Neurosciences, (August, 2002) Vol. 25, No. 8, pp. 390-394.
print.
CODEN: TNSCDR. ISSN: 0166-2236.
DT Article
LA English
ED Entered STN: 14 Aug 2002
Last Updated on STN: 14 Aug 2002

=> index bioscience patents

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FILE 'ENCOMPPAT2' ACCESS NOT AUTHORIZED

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COST IN U.S. DOLLARS		
FULL ESTIMATED COST	39.04	56.19

INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, AQUASCI, BIOBUSINESS, BIOCOMMERCE, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA, CANCERLIT, CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB, CROPU, DISSABS, DDFB, DDFU, DGENE, DRUGB, DRUGMONOG2, ...' ENTERED AT 11:30:07 ON 27 JUL 2004

92 FILES IN THE FILE LIST IN STNINDEX

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=> amplification and centrifuging and heating and "misfolded proteins"

1 FILE DGENE
25 FILES SEARCHED...
1 FILE IFIPAT
50 FILES SEARCHED...
9 FILE USPATFULL
68 FILES SEARCHED...
3 FILE PCTFULL
87 FILES SEARCHED...

4 FILES HAVE ONE OR MORE ANSWERS, 92 FILES SEARCHED IN STNINDEX

L6 QUE AMPLIFICATION AND CENTRIFUGING AND HEATING AND "MISFOLDED PROTEINS"

=> file DGENE IFIPAT USPATFULL PCTFULL
COST IN U.S. DOLLARS

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	ENTRY	SESSION
FULL ESTIMATED COST	3.99	60.18

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=> amplification and centrifuging and heating and "misfolded proteins"

L7 14 AMPLIFICATION AND CENTRIFUGING AND HEATING AND "MISFOLDED PROTEI
NS"

=> d di 1-14

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REENTER DISPLAY FORMAT FOR ALL FILES (FILEDEFAULT):d ti 1-14

'D' IS NOT A VALID FORMAT

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L7 ANSWER 1 OF 14 DGENE COPYRIGHT 2004 THOMSON DERWENT on STN

TI Cyclically amplifying proteons in a biological sample for detecting conformational or congophilic disorders by **centrifuging** the sample until a supernatant is formed, dividing the supernatant into subsamples and **heating** a subsample.

L7 ANSWER 2 OF 14 IFIPAT COPYRIGHT 2004 IFI on STN

TI METHOD OF ISOLATION AND SELF-ASSEMBLY OF SMALL PROTEIN PARTICLES FROM BLOOD AND OTHER BIOLOGICAL MATERIALS

L7 ANSWER 3 OF 14 USPATFULL on STN

TI Method of isolation and self-assembly of small protein particles from blood and other biological materials

L7 ANSWER 4 OF 14 USPATFULL on STN

TI Yeast cell surface display of proteins and uses thereof

L7 ANSWER 5 OF 14 USPATFULL on STN

TI Yeast cell surface display of proteins and uses thereof

L7 ANSWER 6 OF 14 USPATFULL on STN

TI Nucleic acid sequences encoding type III tenebrio antifreeze proteins and method for assaying activity

L7 ANSWER 7 OF 14 USPATFULL on STN

TI Nucleic acid sequences encoding type III tenebrio antifreeze proteins and method for assaying activity

L7 ANSWER 8 OF 14 USPATFULL on STN

TI Yeast cell surface display of proteins and uses thereof

L7 ANSWER 9 OF 14 USPATFULL on STN

TI Yeast cell surface display of proteins and uses thereof

L7 ANSWER 10 OF 14 USPATFULL on STN

TI Yeast cell surface display of proteins and uses thereof

L7 ANSWER 11 OF 14 USPATFULL on STN

TI Production of insulin-like growth factor-1 in methylotrophic yeast cells

L7 ANSWER 12 OF 14 PCTFULL COPYRIGHT 2004 Univentio on STN
TIENT METHOD OF ISOLATION AND SELF-ASSEMBLY OF SMALL PROTEIN PARTICLES FROM
BLOOD AND OTHER BIOLOGICAL MATERIALS
TIFR METHODES D'ISOLEMENT ET D'AUTO-ASSEMBLAGE DE PETITES PARTICULES
PROTEINIQUES PRESENTES DANS DU SANG OU DANS D'AUTRES MATIERES
BIOLOGIQUES

L7 ANSWER 13 OF 14 PCTFULL COPYRIGHT 2004 Univentio on STN
TIENT NUCLEIC ACID SEQUENCES ENCODING TYPE III TENEBRIO ANTIFREEZE PROTEINS
AND METHOD FOR ASSAYING ACTIVITY
TIFR SEQUENCES D'ACIDES NUCLEIQUES CODANT POUR DES PROTEINES ANTIGEL DE
TENEBRION TYPE III ET PROCEDE D'ANALYSE ASSOCIE

L7 ANSWER 14 OF 14 PCTFULL COPYRIGHT 2004 Univentio on STN
TIENT YEAST CELL SURFACE DISPLAY OF PROTEINS AND USES THEREOF
TIFR PROTEINES POUR LA PRESENTATION DE LA SURFACE D'UNE CELLULE DE LEVURE ET
LEURS UTILISATIONS

=> d ab bib 4, 6, 7

L7 ANSWER 4 OF 14 USPATFULL on STN
AB The present invention provides a genetic method for tethering
polypeptides to the yeast cell wall in a form accessible for binding to
macromolecules. Combining this method with fluorescence-activated cell
sorting provides a means of selecting proteins with increased or
decreased affinity for another molecule, altered specificity, or
conditional binding. Also provided is a method for genetic fusion of the
N terminus of a polypeptide of interest to the C-terminus of the yeast
Aga2p cell wall protein. The outer wall of each yeast cell can display
approximately 10.sup.4 protein agglutinins. The native agglutinins serve
as specific adhesion contacts to fuse yeast cells of opposite mating
type during mating. In effect, yeast has evolved a platform for
protein-protein binding without steric hindrance, from cell wall
components. As one embodiment, attaching an scFv antibody fragment to
the Aga2p agglutinin effectively mimics the cell surface display of
antibodies by B cells in the immune system for affinity maturation in
vivo. As another embodiment, T cell receptor mutants can be isolated by
this method that are efficiently displayed on the yeast cell surface,
providing a means of altering T cell receptor binding affinity and
specificity by library screening.

AN 2004:53289 USPATFULL
TI Yeast cell surface display of proteins and uses thereof
IN Wittrup, K. Dane, Chestnut Hill, MA, United States
Kranz, David M., Champaign, IL, United States
Kieke, Michele, Urbana, IL, United States
Boder, Eric T., Media, PA, United States
PA Board of Trustees of the University of Illinois, Urbana, IL, United
States (U.S. corporation)
PI US 6699658 B1 20040302
AI US 1998-9388 19980120 (9)
RLI Continuation-in-part of Ser. No. US 1997-866398, filed on 30 May 1997,
now abandoned
PRAI US 1996-18741P 19960531 (60)
DT Utility
FS GRANTED
EXNAM Primary Examiner: Guzo, David
LREP Greenlee, Winner and Sullivan, P.C.

CLMN Number of Claims: 42
ECL Exemplary Claim: 1
DRWN 36 Drawing Figure(s); 27 Drawing Page(s)
LN.CNT 2835
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 6 OF 14 USPATFULL on STN
AB Thermal hysteresis proteins and their nucleotide sequences derived from the Tenebrionoidea Superfamily which lower the freezing point of a solution without effecting the melting point. Related methods for preparing said proteins and for providing antifreeze or recrystallization inhibition properties to a subject formulation.
AN 2002:307900 USPATFULL
TI Nucleic acid sequences encoding type III tenebrio antifreeze proteins and method for assaying activity
IN Horwath, Kathleen L., Endwell, NY, UNITED STATES
Easton, Christopher M., Ithaca, NY, UNITED STATES
PI US 2002173024 A1 20021121
AI US 2001-876796 A1 20010607 (9)
PRAI US 2000-210446P 20000608 (60)
DT Utility
FS APPLICATION
LREP Mark Levy, SALZMAN & LEVY, Ste. 902, 19 Chenango St., Binghamton, NY, 13901
CLMN Number of Claims: 40
ECL Exemplary Claim: 1
DRWN 131 Drawing Page(s)
LN.CNT 10082
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 7 OF 14 USPATFULL on STN
AB A recrystallization inhibition method for determining the presence, relative concentration, and/or activity of thermal hysteresis proteins comprising: providing a proteinaceous composition in a solvent to form a test solution; flash freezing said solution; raising the temperature of the frozen solution to an appropriate annealing temperature that allows for a partial melt, while limiting heterogeneity in ice grain sizes within said solution; maintaining said frozen solution at the annealing temperature for a length of time sufficient to allow for recrystallization; monitoring the ice crystal grain size changes over time; and determining the presence of functional thermal hysteresis proteins in said solution given the retention of significantly smaller ice crystal grain sizes relative to at least one control solution.
AN 2002:307828 USPATFULL
TI Nucleic acid sequences encoding type III tenebrio antifreeze proteins and method for assaying activity
IN Horwath, Kathleen L., Endwell, NY, UNITED STATES
Meyers, Kevin L., Trumansburg, NY, UNITED STATES
PI US 2002172951 A1 20021121
AI US 2001-876348 A1 20010607 (9)
PRAI US 2000-210446P 20000608 (60)
DT Utility
FS APPLICATION
LREP Mark Levy, SALZMAN & LEVY, Ste. 902, 19 Chenango St., Binghamton, NY, 13901
CLMN Number of Claims: 34
ECL Exemplary Claim: 1
DRWN 131 Drawing Page(s)
LN.CNT 10121